Biomonitoring Quality Assurance Support Program

Analytical Method Report

Environmental Phenols

Laboratory Information

1 State Report:

2 Results for Method:

Sample Preparation Information:

- 3 Does your method use automation:
- 4 Does your sample preparation method include:
- 5 Solid phase extraction platform:
 Additional information on sample prep procedure:
- 6 What is the method sample volume size:
- 7 Name of enzyme:

Enzyme Vendor
Enzyme Concentration
Amount
How long do you incubate the samples
Temperature during incubations

HPLC Configuration:

- 8 Instrument manufacturer
- 9 What is the flow rate:
- 10 What is the method run time:
- 11 What is the sample injection volume:
- 12 Column name and Manufacturer
- 13 Column dimensions
- 14 Elution Type:
- 15 Mobile Phase A Composition
- 16 Mobile Phase B Composition

Mass Spectrometer Configuration

Have you optimized the MS Parameters for your method? (Analytes, 17 Precursor and Product Ions, Collision Energy)

18 What is the ionization mode:

²⁰ Please complete the table for each analytes LOD, precursor and product ion transitions:

Analytical and Internal Standards Please complete the table for metabolite standards:

How do you prepare your standards	25
How many points are in the calibration curve	27
Is the calibration curve weighted	28
What integration software do you use	29

Additional Method Questions

30 Which proficiency testing programs do you participate in?

what is the average number of samples analyzed per month for this 31 method?

32 Have you checked the accuracy of the method using NIST SRMs?

33 What volume of sample is required for BQASP Analysis?

Please provide a screenshot of your results chromatography:

CDC estimates the average public reporting burden for this collection of information instructions, searching existing data/information sources, gatheric completing and reviewing the collection of information. An agency may not to a collection of information unless it displays a currently valid OMB Control any other aspect of this collection of information, including suggestions for Review Office, 1600 Clifton Road NE, MS D-74, Atlanta, Georgia 30333; ATTN

Form Approved OMB No. 0920-xxxx Exp. Date xx/xx/20xx

Select State	_
Select Method	_
Yes or No Select SPE N/A	_ -
Type description here	- -
	Degrees Celsius
Select	_Type Other Here - -
Select	- - - Describe composition for Mobile Phase A
	Describe composition for Mobile Phase B

Yes or No Select Mode

Analyte	LOD
Example: MCPP	0.4 ng/ml
13C4-MCPP	

	•
Analytical and Internal Standard	Vendor
Phthalate Metabolites	Cambridge Isotope Laboratory
Select	
	weighted curve: No weighting, 1/X, 1/X^2,Other
Select	Please type other programs here

Right click in the textbox Click format shape Click Fill Options Select picture from your saved file ormation as 45 minutes per response, including the time for ng and maintaining the data/information needed, and conduct or sponsor, and a person is not required to respond of Number. Send comments regarding this burden estimate or reducing this burden to CDC/ATSDR Information Collection N: PRA (0920-xxxx).

Calibration Range	Precursor Ion (mass)	Product Ion (mass)
0.035 - 350 ng/ml	251	103
	225	103

Purity

?

ENVIRONMENTAL PHENOLS

PT Event ID: 201902EPSU

Participant: Analyst: Reviewer: Units of Result:

Sample ID	<u>Analyte</u>
201902001EPSU	2,4-dichlorophenol
201902001EPSU	2,5-dichlorophenol
201902001EPSU	Butyl paraben
201902001EPSU	Benzophenone-3
201902001EPSU	Bisphenol A
201902001EPSU	Bisphenol S
201902001EPSU	Bisphenol F
201902001EPSU	Ethyl-paraben
201902001EPSU	Methyl-paraben
201902001EPSU	Propyl-paraben
201902001EPSU	Triclocarban
201902001EPSU	Triclosan

<u>Sample ID</u>	<u>Analyte</u>
201902002EPSU	2,4-dichlorophenol
201902002EPSU	2,5-dichlorophenol
201902002EPSU	Butyl paraben
201902002EPSU	Benzophenone-3
201902002EPSU	Bisphenol A
201902002EPSU	Bisphenol S
201902002EPSU	Bisphenol F
201902002EPSU	Ethyl-paraben
201902002EPSU	Methyl-paraben
201902002EPSU	Propyl-paraben
201902002EPSU	Triclocarbon
201902002EPSU	Triclosan

By submitting this form, we attest that the results reported were produced in this laber analysis of proficiency testing samples that were introduced into the routine workflow laboratory and analyzed using protocols and procedures with the same frequency rou patient specimens.

We further attest that the laboratory did not discuss or engage in any communication

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Reported Value

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